CRF Problem Report



The Scientific and Technical Information Center (STIC) experienced problem when processing the following computer readable form (CRF):

2 9 2001 Application Serial Number 09/772, 134

Filing Date:

Date Processed by STIC: 2-9-01

STIC Contact: Mark Spencer, 703-308-4212

Nature of Problem:

The CRF (was):

(circle one) Damaged or Unreadable (for Unreadable, see attached)

Blank (no files on CRF) (see attached)

Empty file (filename present, but no bytes in file) (see attached)

Virus-infected. Virus name: _____ The STIC will not process the CRF.

Not saved in ASCII text

Sequence Listing was embedded in the file. According to Sequence Rules, submitted file should only be the Sequence Listing.

Did not contain a Sequence Listing. (see attached sample)

V Other: Non-valid format, Example of correct formet attached

PLEASE USE THE CHECKER VERSION 3.0 PROGRAM TO REDUCE ERRORS. SEE BELOW FOR DETAILS:

Checker Version 3.0

The Checker Version 3.0 application is a state-of the-art Windows based software program employing a logical and intuitive user-interface to check whether a sequence listing is in compliance with format and content rules. Checker Version 3.0 works for sequence listings generated for the original version of 37 CFR §§1.821 - 1.825 effective October 1, 1990 (old rules) and the revised version (new rules) effective July 1, 1998 as well as World Intellectual Property Organization (WIPO) Standard ST.25.

Checker Version 3.0 replaces the previous DOS-based version of Checker, and is Y2Kcompliant. Checker allows public users to check sequence listings in Computer Readable form (CRF) before submitting them to the United States Patent and Trademark Office (USPTO). Use of Checker prior to filing the sequence listing is expected to result in fewer errored sequence listings, thus saving time and money.

Checker Version 3.0 can be down loaded from the USPTO website at the following address: http://www.uspto.gov/web/offices/pac/checker

Appendix A To Subpart C to Part 1-Sample Sequence Listing



<110> Smith, John

Smith, Jane

<120> Example of a Sequence Listing

<130> 01-00001

<140> US 08/999,999

<141> 1998-02-28

<150> EP 91000000

<151> 1997-12-31

<170> PatentIn ver. 2.0

<210> 1

<211> 403

<212> DNA

<213> Paramecium aurelía

<220>

<221> CDS

<222> 341..394

<300>

<301> Doe, Richard

<302> Isolation and Characterization of a Gene Encoding a

Protease from Paramecium sp.

<303> Journal of Fictional Genes

<305> 4

<306> 1 - 7

<307> 1988-06-20

<400> 1

ctactctact ctactctcat ctactatett ctttggatet ctgagtetge ctgagtggta

ctcttgagtc ctggagatct ctcctctcac atgtgatcgt cgagactgac cgatagatcg 120

ctgactgact ctgagatagt cgagcccgta cgagacccgt cgagggtgac agagagtggg 180

cgcgtgcgcg cagagcgccg cgccggtgcg cgcgcgagtg cgcggtgggc cgcgcgaggg 240

ctttcgcggc agcggcggcg ctttccggcg cgcgcccgtc cgcccctaga cctgagaggt

cttctcttcc ctcctctca ctagagaggt ctatatatac atg gtt tca atg ttc

Met Val Ser Met Phe

age ttg tet tte aaa tgg cet gga ttt tgt ttg ttt gtt tgtttgete

403

Ser Leu Ser Phe Lys Trp Pro Gly Phe Cys Leu Phe Val

10

15

<210> 2

<211> 18

<212> PRT

<213> Paramecium aurelia

<400> 2

bl= 630 No.; 104 / Monday, June 1, 199 ules and

Met Val Ser Met Phe Ser Leu Ser Phe Lys Trp Pro Gly Phe dys Leu

5

10

Phe Val

1

21305		Mark with a contract to the contract to		Street, and a st	, ,	15 company
<110> ,	• • • • • • • • • • • • • • • • • • • •	Applicant	···· [·	Preferably max. of 10 names; one name	line;	M
	• •		- 1	or initials:	and/	
<120>	••••••	Title of Intension		Personal file releases		1
<130>	•••••••	File Reference		Personal file reference		M. M when filed prior to assignment of appl. nu. ber.
<140>		Current Application	i i		••••••	ber. hed phor to assignment of appl. nu.
	••••••	Number.	- 13	Specily as: US 07/999,999 or PCT/US96/999	99	M. il available
i <141>		Current Filing Date	١.			
\$ 150>		Prior Apolication Num		Specify as: yyyy-mm-dd Specify as: US 07/999,999 or PCT/US96/999		M, if available.
		ber.		the second secon	1	M, if applicable include priority documents und
(131>		Prior Application Filin Date.	g S	pecily as: yyyy-mm-dd	-	35.USC:119 and 120. M. If applicable.
<160>		Number of SEQ ID N		•		man approache.
<170>		Software		Count Includes total number of SEQ ID NOs	[M
210	- 1			lame of software used to create the Sequel Listing.	nce	O.
≪10>		SEQ ID NO:1:	R	esponse shall be an integer representing	the	М.
. <211>	1	المحتدد عليه	~ ~ .	i serien un suomu.	- 1	
14.		Length	R	espond with an integer expressing the numi	ber	M. · • •
700				of bases or amino acid residues.		<u> </u>
Númeric Id	en-	Definition	- 1	Comments and format	-T	No.
				Solutions and ionial		Mandatory (M) or optional (O).
<212>	1	Гуре	wı	hether presented sequence molecule is DN	A	Λ.
			1 8	RNA, or PRT (protein). If a nucleotide s	c.	To a mandarp
			1 (quence contains both DNA and RNA Ira	n- 1	
	- 1		l ü	ments, the type shall be "DNA." In addition he combined DNA/RNA molecule shall be fu	n.	and the same of th
	į		j u	her described in the <220> to <223> feature	re	
<213×		Organism	3	ection,	Į.	
, . · · · · · · · · · · · · · · · · · ·				entific name, i.e. Genus/ species, Unknown o	or M	
		•	هُ. ا	rtificial Sequence. In addition, the "Unknown ar "Artificial Sequence" organisms shall be ful	ו"ן	$g_{a}(x) = e^{-x^{2}}$
	-	·	l n	her described in the <220> to <223> featur	e	*
<220>	Fo	calure	5€	ection,	- 1	•
				ve blank after <220>. <221-223> provide fo description of points of biological significance	r M,	under the following conditions: if "n," "Xaa,"
			In	the sequence	יוי	or a mounicu of unusual Leaming acid as mad:
			1	•		fied base was used in a sequence; if ORGA- NISM is "Artificial Sequence" or "Unknown; if
<221>	_{Na}	ime/Kev	0	tale and the second of the sec		HOICECULE IS COMPUTED DIVATED A.
			. Prov	ride appropriate Identifier for feature, pref-	· M,	under the following conditions: if "a " "van "
-000·	1.		pe	ably from WIPO Standard ST.25 (1998), Ap- endix 2, Tables 5 and 6.	1 6	o income of unusual traming a sid or mind.
<222>	Lo	cation	Spec	city location within sequence; where appro-	М.	under the following conditions: if "a " "Year"
			pna	ate state number of first and last bases/ nino acids in feature.	0	" a mounted of unusual L-amino acid or mod:
<223>	. Oth	er Information	Other	ino acids in leature. r relevant information; four lines maximum	, ,,	ed base was used in a sequence
				and maximum	M.	under the following conditions: if "a " "vaa "
			1		1110	r a modified or unusual L-amino acid or modi- ed base was used in a sequence; if ORGA-
	1				1 14	ISM IS ANIIIQAI Sequence" or "Hekaniii" ir
<300>	Pub	lication Information	Leave	blank after <300>		olecule is combined DNA/RNA.
<301>	Auth	nors	Prefer	rably max of ten named authors of publica-	0. 0.	
	1		lion	: specify one name per line; preferable for-		
<302>		***************************************	mal	: Surname, Other Names and/or Initials.		``
<303>	Jour	**********************			O. O.	··
<304> <305>	Volu	me			O.	
<306>	Page	28	•••••		Ö.	įi
<307>	Date		Journa	I date on which data published	0.	. `
			уууу	date on which data published; specify as mm-dd, MMM-yyyy or Season-yyyy.	Ο.	•
<308>	Data	base Accession	Access	sion number assigned by database include	Ο.	
<309>		moer	ing d	Jalabase name.		
			dd o	f entry in database; specify as yyyy-mm- r MMM-yyyy.	Ο.	* -
<310>		nt Document Num-	Docum	ent number; for patent-type citations only	Ο.	
<311>	ber Paten	·	Spec	ily as, for example, US 07/999 999	J .	
1	aten	t Filing Date	Docume	ent filing date, for patent-type citations to	Ο.	. •
<312>	Public	ation Date	Docum	specify as yyyy-mm-dd. ent publication date, for patent-type cita-	_	
j			TIONS	only; specify as vvvv-mm-dd	Э.	
<400>	Secur	ant Residuesz	-ном ((position) TO (position)) .	
	Jeque		שני שפ	NO should follow the numeric identifier L	A .	
_			actual	should appear on the line preceding the sequence.		